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Listing of Claims

The listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently Amended) A compound of formula (I):

$$\begin{array}{c|c}
R^2 \\
Q^2 \\
R^3 \\
Q^3 \\
Q^1
\end{array}$$

$$\begin{array}{c|c}
Q^1 \\
R^b \\
R^1
\end{array}$$

wherein

Y1 is CH or N;

Q1 is selected from the group consisting of

- (1) -OH, and
- $(2) NH_2;$

Q² and Q³ independently selected from the group consisting of

- (1) hydrogen, and
- (2) halogen;

Ra is selected from the group consisting of

- (1) hydrogen,
- (2) -C₁₋₁₀ alkyl, wherein said alkyl is unsubstituted or substituted with one or more fluoro, and
- (3) -C3-8 cycloalkyl;

Rb is selected from the group consisting of

- (1) hydrogen,
- $(2) C_{1-10}$ alkyl,
- (3) -C₁₋₃ alkyl-aryl, wherein said aryl is selected from the group consisting of phenyl and naphthyl,
- (4) -C3-8 cycloalkyl,

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wherein said cycloalkyl, alkyl and aryl are unsubstituted or substituted with one or more

- (a) halo,
- (b) -OH,
- (c) -CN,
- (d) -O-C₁₋₁₀ alkyl,
- (5) $-(CH_2)_n$ -NRCRd wherein RC and Rd are selected from the group consisting of hydrogen and C_{1-10} alkyl, and n is 2, 3 or 4, and
- (6) -(CH₂)_n,-O-Re, wherein Re is selected from the group consisting of
 - (a) C₁₋₁₀ alkyl,
 - (b) -C₀₋₃ alkyl-aryl, wherein said aryl is selected from the group consisting of phenyl and naphthyl,

wherein said alkyl and aryl are unsubstituted or substituted with one or more

- (i) halo,
- (ii) -OH,
- (iii) -CN,
- (iv) $-O-C_{1-10}$ alkyl,

and n' is 1, 2, 3 or 4;

m is 1 or 2;

- R¹ is (1) aryl selected from the group consisting of phenyl and napthyl, or
 - (2) heteroaryl selected from the group consisting of pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyridyl, pyridyl, tetrazolyl, furanyl, imidazolyl, triazinyl, pyranyl, thiazolyl, thienyl, thiophenyl, triazolyl, oxazolyl, isoxazolyl, thiazolyl, oxadiazolyl, indolyl, quinolinyl, isoquinolinyl, benzimidazolyl and benzoxazolyl,
 - (3) -C₁₋₁₀ alkyl, and
 - (4) -C₃₋₈ cycloalkyl,

wherein said aryl, heteroaryl, alkyl and cycloalkyl is unsubstituted or substituted with one or more

- (a) halo,
- (b) -OH,
- (c) -CN,
- (d) $-O-C_{1-10}$ alkyl,

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- (e) $-C_{1-10}$ alkyl,
- (f) -C3-8 cycloalkyl,
- (g) aryl selected from the group consisting of phenyl and napthyl, or
- (h) heteroaryl selected from the group consisting of pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolyl, tetrazolyl, furanyl, imidazolyl, triazinyl, pyranyl, thiazolyl, thienyl, thiophenyl, triazolyl, oxazolyl, isoxazolyl, thiazolyl, oxadiazolyl, indolyl, quinolinyl, isoquinolinyl, benzimidazolyl and benzoxazolyl;

R² is selected from the group consisting of:

- (1) $(R^4-SO_2)N(R^7)$ -, wherein R^4 is
 - (a) -C₁₋₁₀ alkyl,
 - (b) -C₃₋₈ cycloalkyl,

wherein said alkyl and cycloalkyl is unsubstituted or substituted with one or more

- (i) halo,
- (ii) -OH,
- (iii) -CN,
- (iv) $-O-C_{1-10}$ alkyl,
- (v) $-C_{1-10}$ alkyl,
- (vi) -C3-8 cycloalkyl,
- (vii) aryl selected from the group consisting of phenyl and napthyl, or (viii) heteroaryl selected from the group consisting of pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolyl, tetrazolyl, furanyl, imidazolyl, triazinyl, pyranyl, thiazolyl, thienyl, thiophenyl, triazolyl, oxazolyl, isoxazolyl, thiazolyl, oxadiazolyl, indolyl, quinolinyl, isoquinolinyl, benzimidazolyl and benzoxazolyl;

and said aryl and heteroaryl is unsubstituted or substituted with one or more

- (A) halo,
- (B) -OH,
- (C) -CN,
- (D) $-O-C_{1-10}$ alkyl,
- (E) -C₃₋₈ cycloalkyl, or
- (F) -C₁₋₁₀ alkyl,

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(c) heteroaryl selected from the group consisting of pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolyl, tetrazolyl, furanyl, imidazolyl, triazinyl, pyranyl, thiazolyl, thienyl, thiophenyl, triazolyl, oxazolyl, isoxazolyl, thiazolyl, oxadiazolyl, indolyl, quinolinyl, isoquinolinyl, benzimidazolyl and benzoxazolyl,

wherein said heteroaryl is unsubstituted or substituted with one or more

- (i) halo,
- (ii) -OH,
- (iii) -CN,
- (iv) $-O-C_{1-10}$ alkyl,
- (v) -C3-8 cycloalkyl, or
- (vi) -C1-10 alkyl,

(d) – $(CH_2)_X$ -NR^fRg wherein R^f and Rg are selected from the group consisting of hydrogen and C_{1-10} alkyl, and x is 0, 1, 2, 3 or 4, or R^f and Rg, together with the nitrogen atom to which they are attached form the group

wherein y is 1 or 2, Y⁵ is -CHR²¹, -O- or NR²¹, wherein R²¹ is selected from the group consisting of;

- (i) hydrogen, and
- (ii) C₁₋₁₀ alkyl,

wherein said alkyl is unsubstituted or substituted with one or more

- (A) halo,
- (B) -OH,
- (C) -CN,
- (D) -O-C1-10 alkyl, or
- (E) -C₃₋₈ cycloalkyl;

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R⁷ is selected from the group consisting of

- (a) hydrogen, and
- (b) $-C_{1-10}$ alkyl,
- (c) aryl selected from the group consisting of phenyl and napthyl, or
- (d) heteroaryl selected from the group consisting of pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolyl, tetrazolyl, furanyl, imidazolyl, triazinyl, pyranyl, thiazolyl, thienyl, thiophenyl, triazolyl, oxazolyl, isoxazolyl, thiazolyl, oxadiazolyl, indolyl, quinolinyl, isoquinolinyl, benzimidazolyl and benzoxazolyl

wherein said alkyl, aryl and heteroaryl is unsubstituted or substituted with one or more

- (i) halo,
- (ii) -OH,
- (iii) -CN,
- (iv) $-O-C_{1-10}$ alkyl,
- (v) -C₃₋₈ cycloalkyl,
- (vi) aryl selected from the group consisting of phenyl and napthyl, or (vii) heteroaryl selected from the group consisting of pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolyl, tetrazolyl, furanyl, imidazolyl, triazinyl, pyranyl, thiazolyl, thienyl, thiophenyl, triazolyl, oxazolyl, isoxazolyl, thiazolyl, oxadiazolyl, indolyl, quinolinyl, isoquinolinyl, benzimidazolyl and benzoxazolyl,

wherein said cycloalkyl, aryl or heteroaryl is unsubstituted or substituted with one or more

- (A) halo,
- (B) -OH,
- (C) -CN,
- (D) $-O-C_{1-10}$ alkyl,
- (E) -C₃₋₈ cycloalkyl, or
- (F) aryl selected from the group consisting of phenyl and napthyl;
- (e) –(CH₂)_y'–NR^hRⁱ wherein R^h and Rⁱ are selected from the group consisting of hydrogen and C₁₋₁₀ alkyl, and y' is 1, 2, 3 or 4, or or R^h and Rⁱ, together with the nitrogen atom to which they are attached from the group

wherein y' is 1 or 2, Y6 is -CHR²², -O- or NR²², wherein R²² is selected from the group consisting of;

- (i) hydrogen, and
- (ii) C₁₋₁₀ alkyl,

wherein said alkyl is unsubstituted or substituted with one or more

- (A) halo,
- (B) -OH,
- (C) -CN,
- (D) -O-C₁₋₁₀ alkyl, or
- (E) -C3-8 cycloalkyl,

or R⁴ and R⁷ are linked together to form the group

(a)

wherein z is 1, 2 or 3; or

(b)

wherein z is 1, 2 or 3

(2)

wherein R⁸ is selected from the group consisting of

- (a) -CN,
- (b) hydrogen, and
- (c) tetrazolyl;

(3)

(4)

wherein Y² is -NH=CH- or -CH=NH-;

R³ is selected from the group consisting of

wherein Y3 is CR6c or N;

R⁵ is C₁₋₁₀ alkyl or C₁₋₂ perfluoroalkyl;

R6a, R6b, and R6c are independently selected from the group consisting of:

(1) hydrogen,

(2) halo,

- $(3) C_{1-10}$ alkyl,
- (4) -OH,
- (5) –CN,
- (6) -C3-8 cycloalkyl, and
- (7) -O-C₁₋₁₀ alkyl;

R9 and R10 are independently selected from the group consisting of

- (1) hydrogen,
- (2) -C₁₋₁₀ alkyl, and
- (3) -C3-8 cycloalkyl,

wherein said alkyl and cycloalkyl are unsubstituted or substituted with one or more

- (a) halo,
- (b) -OH,
- (c) -CN,
- (d) -O-C₁₋₁₀ alkyl,
- (e) -C3-8 cycloalkyl, and
- (f) -NRj Rk wherein Rj and Rk are C1-10 alkyl;

or R9 and R10 are joined together with the nitrogen atom to which they are attached to form

$$R^{23}$$
 r^{ν}

wherein w is 1, 2 or 3, and

R²³ is selected from the group consisting of

- (a) hydrogen,
- (b) -C₁₋₁₀ alkyl,
- (c) -C₃₋₈ cycloalkyl,
- (d) $-C_{2-10}$ alkenyl,
- (e) -C2-10 alkynyl,
- (f) -(CH₂)_p-phenyl,

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(g) -(CH₂)_p-heteroaryl, wherein said heteroaryl is selected from the group consisting of pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolyl, tetrazolyl, furanyl, imidazolyl, triazinyl, pyranyl, thiazolyl, thienyl, thiophenyl, triazolyl, oxazolyl, isoxazolyl, thiazolyl, oxadiazolyl, indolyl, quinolinyl, isoquinolinyl, benzimidazolyl and benzoxazolyl,

wherein p is 0 or 1, and

wherein said alkyl, alkenyl, alkynyl, cycloalkyl, phenyl and heteroaryl is unsubstituted or substituted with one or more

- (i) halo,
- (ii) -C₁₋₁₀ alkyl,
- (iii) -OH,
- (iv) -CN,
- (v) -C3-8 cycloalkyl, or
- (vi) -O-C₁₋₁₀ alkyl;

R¹¹ is selected from the group consisting of

- (1) CH -
- (2) –CH₂–,
- (3) O -, and
- $(4) NR^{17} -$

provided that when R¹¹ is -CH- the dotted line forms a bond and when R¹¹ is -CH₂-, -O- or -NR¹⁷- the dotted line is absent;

 R^{17} is hydrogen or C_{1-10} alkyl, wherein said C_{1-10} alkyl is unsubstituted or substituted with one or more

- (a) halo,
- (b) -OH,
- (c) -CN,
- (d) -C3-8 cycloalkyl,
- (e) $-O-C_{1-10}$ alkyl,
- (f) $-(CH_2)_q$ -phenyl, wherein q is 1 or 2, and
- (g) -NR18R19, and

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wherein R18 and R19 are independently selected from the group consisting of

- (i) hydrogen, or
- (ii) C₁₋₁₀ alkyl;

or R18 and R19, together with the nitrogen atom to which they are attached, form the group



wherein q' is 1 or 2, Y⁷ is -CHR²⁴, -O- or NR²⁴, wherein R²⁴ is selected from the group consisting of;

- (a) hydrogen, and
- (b) C₁₋₁₀ alkyl,

wherein said alkyl is unsubstituted or substituted with one or more

- (i) halo,
- (ii) -OH,
- (iii) -CN,
- (iv) $-O-C_{1-10}$ alkyl, or
- (v) -C3-8 cycloalkyl;

R²⁶ is selected from the group consisting of

- (1) hydrogen,
- (2) $-C_{1-3}$ alkyl;

R12 is selected from the group consisting of

- (1) hydrogen,
- (2) -C₁₋₁₀ alkyl, wherein said alkyl is unsubstituted or substituted with one or more
 - (a) halo,
 - (b) -OH,
 - (c) -CN,
 - (d) -C3-8 cycloalkyl,
 - (e) -O-C₁₋₁₀ alkyl, or
 - $(f) -NH_2$

- (3) halo,
- (4) -C₃₋₈ cycloalkyl,
- (5) aryl selected from the group consisting of phenyl and napthyl, and
- (6) heteroaryl selected from the group consisting of pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolyl, tetrazolyl, furanyl, imidazolyl, triazinyl, pyranyl, thiazolyl, thienyl, thiophenyl, triazolyl, oxazolyl, isoxazolyl, thiazolyl, oxadiazolyl, indolyl, quinolinyl, isoquinolinyl, benzimidazolyl and benzoxazolyl,

wherein said aryl and heteroaryl is unsubstituted or substituted with one or more

- (a) halo,
- (b) -OH,
- (c) -CN,
- (d) $-O-C_{1-10}$ alkyl,
- (e) -C3-8 cycloalkyl, or
- (f) $-C_{1-10}$ alkyl;

R¹³ is selected from the group consisting of

- (1) hydrogen,
- (2) C₁₋₁₀ alkyl, and
- (3) -C₃₋₈ cycloalkyl;

wherein said alkyl and cycloalkyl is unsubstituted or substituted with one or more

- (a) halo,
- (b) -OH,
- (c) -CN,
- (d) -C₃₋₈ cycloalkyl,
- (e) -O-C₁₋₁₀ alkyl, and
- (f) $-C_{1-10}$ alkyl;

R¹⁴ is selected from the group consisting of

- (1) -C₁₋₁₀ alkyl, and
- (2) -C3-8 cycloalkyl;

wherein said alkyl and cycloalkyl is unsubstituted or substituted with one or more

- (a) halo,
- (b) -OH,
- (c) -CN,

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(d) -C3-8 cycloalkyl,

- (e) -O-C₁₋₁₀ alkyl, or
- (f) $-C_{1-10}$ alkyl;
- (3) -(CH₂)_V-NR¹⁵R¹⁶, wherein v is 2, 3 or 4, and wherein R¹⁵ and R¹⁶ are independently selected from the group consisting of
 - a) hydrogen, or
 - b) C₁₋₁₀ alkyl, wherein said C₁₋₁₀ alkyl is unsubstituted or substituted with one or more
 - (i) halo,
 - (ii) -OH,
 - (iii) -CN,
 - (iv) -C3-8 cycloalkyl, or
 - (v) $-O-C_{1-10}$ alkyl;

or R^{15} and R^{16} , together with the nitrogen atom to which they are attached, form the group



wherein s is 1 or 2, Y^4 is -CHR²⁴-, -O- or -NR²⁴-, wherein R²⁴ is selected from the group consisting of

- (i) hydrogen, and
- (ii) C₁₋₁₀ alkyl,

wherein said alkyl is unsubstituted or substituted with one or more

- (A) halo,
- (B) -OH,
- (C) -CN,
- (D) -O-C₁₋₁₀ alkyl, or
- (E) -C3-8 cycloalkyl,
- 4) -(CH₂)_r-phenyl, wherein r is 1, 2, 3, or 4, and

wherein said phenyl is unsubstituted or substituted with one or more

- (a) halo,
- (b) -OH,

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(c) -CN,

- (d) -O-C₁₋₁₀ alkyl,
- (e) -C₃₋₈ cycloalkyl, or
- (f) -C₁₋₁₀ alkyl;

or R13 and R14, together with the nitrogen atom to which they are attached, form the group



wherein u is 1 or 2, Y⁸ is -CHR²⁵-, -O- or -NR²⁵-, wherein R²⁵ is selected from the group consisting of

- (a) hydrogen,
- (b) C₁₋₁₀ alkyl,
- (c) -(CH2)t-phenyl,
- (d) -(CH₂)_t-heteroaryl, wherein said heteroaryl is selected from the group consisting of pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolyl, tetrazolyl, furanyl, imidazolyl, triazinyl, pyranyl, thiazolyl, thienyl, thiophenyl, triazolyl, oxazolyl, isoxazolyl, thiazolyl, oxadiazolyl, indolyl, quinolinyl, isoquinolinyl, benzimidazolyl and benzoxazolyl,

wherein t is 0 or 1, and

wherein said alkyl, phenyl and heteroaryl is unsubstituted or substituted with one or more

- (i) halo,
- (ii) $-C_{1-10}$ alkyl,
- (iii) -OH,
- (iv) -CN,
- (v) -C3-8 cycloalkyl, or
- (vi) $-O-C_{1-10}$ alkyl;

or a and pharmaceutically acceptable salt salts thereof.

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2. (Currently Amended) The compound of Claim 1, or a pharmaceutically acceptable salt thereof, wherein R^a and R^b are both hydrogen.

- 3. (Currently Amended) The compound of Claim 1, or a pharmaceutically acceptable salt thereof, wherein Ra is hydrogen and Rb is C1-10 alkyl.
- 4. (Currently Amended) The compound of Claim 1, or a pharmaceutically acceptable salt thereof, wherein m is 1 and R¹ is selected from the group consisting of
 - (1) phenyl, unsubstituted or substituted in one or two positions with halo; and
 - (2) thienyl.
- 5. (Currently Amended) The compound of Claim 1, or a pharmaceutically acceptable salt thereof, wherein R² is (R⁴-SO₂)N(R⁷)-.
- 6. (Currently Amended) The compound of Claim 5, or a pharmaceutically acceptable salt thereof, wherein R4 and R7 are each C_{1-6alkyl}.
- 7. (Currently Amended) The compound of Claim 1, or a pharmaceutically acceptable salt thereof, wherein R³ is (1)

$$\begin{array}{c|c}
R^{6a} \\
R^{6b} & \downarrow \downarrow \\
 & \downarrow \downarrow \downarrow 3 \\
R^5 & O
\end{array}$$

wherein Y³ is CHR^{6c}, R⁵ is methyl, R^{6a} and R^{6c} are hydrogen and R^{6b} is fluoro.

8. (Currently Amended) The compound of Claim 1, or a pharmaceutically acceptable salt thereof, wherein R³ is (1)

 Y^3 is N, R^5 is C_{1-2} perfluoroalkyl, and R^{6a} and R^{6b} are hydrogen.

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9. (Currently Amended) The compound of Claim 1, or a pharmaceutically acceptable salt thereof, wherein R³ is (2)

and R⁹ and R¹⁰ are each unsubstituted C₁₋₁₀ alkyl, or R⁹ and R¹⁰ are joined together with the nitrogen atom to which they are attached to form attached to form

$$R^{23}$$
 S^{3}

wherein w is 1;

R²³ is -(CH₂)_p-phenyl or -(CH₂)_p-heteroaryl, wherein said heteroaryl is selected from the group consisting of pyrazinyl, pyrazolyl, pyridazinyl, pyridyl, pyrimidinyl, pyrrolyl, tetrazolyl, furanyl, imidazolyl, triazinyl, pyranyl, thiazolyl, thienyl, thiophenyl, triazolyl, oxazolyl, isoxazolyl, thiazolyl, oxadiazolyl, indolyl, quinolinyl, isoquinolinyl, benzimidazolyl and benzoxazolyl,

wherein the phenyl and heteroaryl are unsubstituted or substituted with one or more chloro, and p is 0.

10. (Currently Amended) The compound of Claim 1, or a pharmaceutically acceptable salt thereof, wherein R^3 is (3)

$$R^{12}$$
 R^{11}

R11 is NR17 wherein R17 is hydrogen or C1-3 alkyl, and R12 is hydrogen or methyl.

11. (Currently Amended) The compound of Claim 1, or a pharmaceutically acceptable salt thereof, wherein R³ is (4)

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 R^{13} is hydrogen and R^{14} is $-(CH_2)_{v}$ -NR¹⁵R¹⁶ wherein v is 2 and R^{15} and R^{16} are each C_{1-10} alkyl, which is unsubstituted or substituted with -OH, -CN or -OCH₃.

12. (Currently Amended) The compound of Claim 1, or a pharmaceutically acceptable salt thereof, wherein R³ is (4)

wherein R¹³ and R¹⁴, together with the nitrogen atom to which they are attached, form the group

wherein u is 1 or 2, Y8 is -CHR²⁵-, -O- or -NR²⁵-.

13. (Currently Amended) The compound of Claim 1 which is a compound of formula (II)

wherein Q¹, Q², Q³, R^a, R^b, R¹, R², R¹², R¹⁷, R²⁶ and m are as defined in Claim 1, or a and pharmaceutically acceptable salt salts thereof.

14. (Currently Amended) The compound of Claim 1 which is a compound of formula (III)

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$$R^{14} \longrightarrow N \longrightarrow Q^{2} \longrightarrow R^{a} \longrightarrow Q^{1}$$

$$R^{14} \longrightarrow R^{13} \longrightarrow Q^{3} \longrightarrow R^{1}$$

$$R^{1} \longrightarrow R^{1}$$

$$R^{1} \longrightarrow R^{1}$$

$$R^{1} \longrightarrow R^{1}$$

wherein Q1, Q2, Q3, Ra, Rb, R1, R2, R13, R14 and m are defined as in Claim 1, or a and pharmaceutically acceptable salt salts thereof.

15. (Currently Amended) The compound of Claim 1 which is a compound of formula (IV):

$$\begin{array}{c|c}
R^2 & Q^2 & R^a \\
\hline
Q^3 & Q & Q^1 \\
\hline
IV & R^1
\end{array}$$

wherein Q^1 , Q^2 , Q^3 , R^a , R^b , R^1 , R^2 and m are as defined in Claim 1, and R^3 is (1) or (2) as defined in Claim 1, or a and pharmaceutically acceptable <u>salt</u> salts thereof.

16. (Currently Amended) A compound of claim 1 which is selected from the group consisting of

D H N H N H N H N H N H N H N H N H N H	N H N OH OH
O N H N H N H N H N H N H N H N H N H N	O S H N H O H F
O O O O O O O O O O O O O O O O O O O	O H N H CI
	N H N H OH OH
Z H O H O H	

-Z -Z - O H Z - I - Z - O H Z - I - Z - O H Z - I - Z - O H Z - I - Z - O H Z - I - Z - O H Z - I - Z - O H Z - I - Z - O H Z - D - Z
N N H ON S
H Z H

	$ \begin{array}{c} $
O N N N N N N N N N N N N N N N N N N N	D H N H ₂

O N N N N N N N N N N N N N N N N N N N	
H N OH F	D D D D D D D D D D D D D D D D D D D
D D D D D D D D D D D D D D D D D D D	N H NH2
NH2 NH2 NH2	NC H NH2 N Ph
NC H NH ₂ N - Ph	NC H OH N Ph

NC H OH N E Ph	NC H NH ₂ N D D D D D D D D D D D D D D D D D D D
NC H NH ₂ N = Ph	NC H NH ₂ N - Ph
F H OH N E Ph	MeO ₂ S. H N N N S S S
MeO ₂ S. H N H NH ₂ Ph	O S O H O H
Z HZ OH	

O N N O H	N N NH OH
CN ONH ONH OH	CN HZ OH F
N, NH N=N O NH O O NH O O O O N	Z IZ O IZ
F CN H OH	CN HN OH OH S
CN CN NOO ON S	Z HZ OH

NC OH OH	O N N O H N O H
T T O H O H N O H	
NH ₂	NH ₂
NH ₂	
	NH ₂ F

NH ₂ F	O NH2 O
NH ₂	
NH ₂	NH ₂
NH ₂	NH ₂

NH ₂	
NH ₂ F	NH ₂ F
F NH2	NH ₂ F
NH ₂ F O N N N N N N N N N N N N N N N N N N	NH ₂ F

NH ₂ F	O O O O O O O O O O O O O O O O O O O
NH ₂ F	NH ₂
O NH ₂ F	O N N N N N N N N N N N N N N N N N N N
ON NH2 F	NH ₂

O D NH2 O NH2 O	O NH ₂ F
	NH ₂ N F
N N N N N N N N N N N N N N N N N N N	NH ₂
NH ₂ NH ₂ NH ₂	NH ₂

NH2 N F F	NH ₂
NH ₂	NH ₂
NH ₂ F _F	NH ₂
NH2 NH2 NH2	NH ₂ NH ₂ NH ₂ F
NH ₂ ONH	NH2. FFF

or a and pharmaceutically acceptable salts salt thereof.

17. (Currently Amended) A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 1, or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier.

- 18. (Canceled)
- 19. (Canceled)
- 20. (Canceled)